



TITLE:

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CITATION:

Tartey, Sarang ...[et al]. Akirin2 is critical for inducing inflammatory genes by bridging $\text{I}\kappa\text{B}-\zeta$ and the SWI/SNF complex.. The EMBO Journal 2014, 33(20): 2332-2348

ISSUE DATE:

2014-10-16

URL:

<http://hdl.handle.net/2433/199669>

RIGHT:

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Akirin2 is critical for inducing inflammatory genes by bridging I κ B- ζ and the SWI/SNF complex

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Running Title: Akirin2 bridges NF- κ B and chromatin modifiers

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Summary

Transcription of inflammatory genes in innate immune cells is coordinately regulated by transcription factors, including NF- κ B, and chromatin modifiers. However, it remains unclear how microbial sensing initiates chromatin remodeling. Here we show that Akirin2, an evolutionarily conserved nuclear protein, bridges NF- κ B and the chromatin remodeling SWI/SNF complex by interacting with BRG1 Associated Factor 60 (BAF60) proteins as well as I κ B- ζ , which forms a complex with the NF- κ Bp50 subunit. These interactions are essential for Toll-like receptor-, RIG-I- and *Listeria*-mediated expression of proinflammatory genes including *Il6* and *Il12b* in macrophages. Consistently, effective clearance of *Listeria* infection required Akirin2. Furthermore, Akirin2 and I κ B- ζ recruitment to the *Il6* promoter depend upon the presence of I κ B- ζ and Akirin2 respectively, for regulation of chromatin remodeling. BAF60 proteins were also essential for the induction of *Il6* in response to LPS stimulation. Collectively, the I κ B- ζ -Akirin2-BAF60 complex physically links the NF- κ B and SWI/SNF complexes in innate immune cell activation.

Introduction

Innate immune cells such as macrophages sense molecular patterns from microorganisms and damaged cells (Beutler, 2009; Medzhitov, 2008). These molecular patterns are recognized by several classes of sensor proteins, such as Toll-like receptors (TLRs), RIG-I-like receptors (RLRs), Nod-like receptors (NLRs) and so on. TLRs and RLRs trigger signaling pathways leading to transcriptional expression of a set of genes involved in inflammation (Takeuchi & Akira, 2010). Systemic inflammation is mediated by the action of proinflammatory cytokines and chemokines such as tumor necrosis factor (TNF), interleukin (IL)-1 β , IL-6, IL-12, and type I IFNs. It has been well documented that transcription factors such as NF- κ B, AP-1 and IFN-regulatory factors (IRFs) are critical for the expression of these inflammatory genes (Honda & Taniguchi, 2006; Oeckinghaus et al, 2011). Moreover, inflammatory gene expression is controlled by various other mechanisms, such as modulation of the decay rate of induced mRNAs, even after nuclear translocation of transcription factors.

Epigenetic regulation of transcription constitutes another such mechanism. Mammalian DNA is tightly packed into chromatin which must be modified by the histone-modifying and ATP-dependent SWI/SNF family of remodeling complexes in order to facilitate gene activation (Medzhitov & Horng, 2009), thereby allowing transcription at specific loci (Cairns, 2009). Several transcription factors can interact with SWI/SNF complexes and recruit them to specific genes (Chi et al, 2004; Clapier & Cairns, 2009; Yudkovsky et al, 1999). For example, IFN- γ restores *IL6* expression by facilitating TLR4-induced recruitment of chromatin remodeling machinery to the *IL6* promoter and promoting *IL6* locus accessibility in tolerized monocytes (Chen & Ivashkiv, 2010). Nucleosome remodeling also appears to contribute to the rapid induction of the p40 subunit of the proinflammatory cytokine interleukin-12 (IL-12) after LPS stimulation in murine macrophages (Weinmann et al, 1999). Nucleosome remodeling requires TLR4 signaling and new protein synthesis but is independent of the NF- κ B subunit c-Rel, which is essential for transcription in macrophages (Weinmann et al, 2001).

The transcriptional induction of mammalian genes by TLRs and other stimuli can be classified in terms of their dependence on SWI/SNF. It was previously proposed that class of

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genes characterized by CpG-island promoters facilitate promiscuous induction from constitutively active chromatin, although another major class consists of non-CpG-island promoters that assemble into stable nucleosomes, resulting in SWI/SNF dependence and a requirement for transcription factors that promote selective nucleosome remodeling (Ramirez-Carrozzi et al, 2009). However, a more recent work suggests that some of CpG-island containing genes also require new protein synthesis and are expressed with slow kinetics (Bhatt et al, 2012). The relationship between chromatin remodeling and transcription factor activation in cytokine gene expression is thus not well understood.

An interesting feature of the innate immune system is the high conservation among species from insects to mammals. Indeed, the role of the Toll protein in innate immunity was discovered through work on *Drosophila*. We previously identified Akirin, a nuclear factor regulating NF- κ B dependent transcription in a functional genome-wide RNA-mediated interference (RNAi) screening of *Drosophila* cell culture to isolate new components of the Imd pathway (Goto et al, 2008). Akirins are found in diverse animal species and possess two conserved regions at their N- and C-termini, the former containing a nuclear localization signal. Akirin has been reported to be crucial for regulating gene expression in a wide range of contexts (Goto et al, 2008; Macqueen & Johnston, 2009; Nowak et al, 2012; Salerno et al, 2009). It has also been established that Akirin has a promyogenic role during muscle regeneration in mice (Marshall et al, 2008). There are 2 Akirin family members in humans and mice. Akirin2, but not Akirin1 is critical for the production of IL-6 in response to TLR stimuli in mouse embryonic fibroblasts (MEFs) in addition to having a role in mouse development (Goto et al, 2008). As Akirin sequences show no obvious DNA- or RNA-binding motifs, they represent a potential link between NF- κ B induced transcription and upstream signaling.

In the present study, we examined the role of mouse Akirin2 in macrophages by generating a conditional allele. Akirin2 is essential for the expression of a set of inflammatory genes including *Il6* downstream of TLRs and RLRs. In addition, mice lacking Akirin2 in macrophages show impaired cytokine production in response to *Listeria* infection and clearance of infecting bacteria *in vivo*. Akirin2-dependent genes tended to exhibit relatively

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few CpG islands in their promoters. These observations motivated us to examine how Akirin2 regulates inflammatory gene expression. Akirin2 was directly recruited to its target gene promoters, and was found to control chromatin remodeling by recruiting BAF60 proteins, components of the SWI/SNF complex. Further, we identified I κ B- ζ as an Akirin2 binding protein via the C-terminal region of Akirin2, and found that I κ B- ζ and the NF- κ B p50 subunit are required for the recruitment of Akirin2 to the *Il6* promoter. Reciprocally, I κ B- ζ was also recruited to the *Il6* promoter in the presence of Akirin2. This study reveals that Akirin2 mediates the physical link between the NF- κ B and SWI/SNF complexes and thereby represents a novel paradigm for providing tissue and target specificity for transcription factor interactions with chromatin remodeling machinery.

Results

Akirin2 Deletion Severely Impairs Proinflammatory Cytokine Production in Macrophages

To investigate the role of Akirin2 in innate immune cells, we specifically ablated Akirin2 in myeloid cells by crossing *Akirin2^{fl/fl}* mice with mice expressing the lysozyme promoter-driven cre recombinase gene (LysM-Cre). Selective deletion of Akirin2 in myeloid cells of *LysM^{Cre/+};Akirin2^{fl/fl}* mice was demonstrated by northern blot and immunoblot analyses (**Supplementary Figure S1A, S1B**). Immunoblot analysis showed a modest decrease in Akirin2 protein expression in macrophages upon LPS stimulation (**Supplementary Figure S1B**). The frequency of T and B cells, macrophages and neutrophils in the spleen, bone marrow (BM) and peritoneal exudate cells (PECs), were not altered between *LysM-Cre⁺;Akirin2^{fl/+}* or *LysM-Cre⁺;Akirin2^{fl/fl}* mice (**Supplementary Figure S2A, S2B, S2C**).

We first examined the responses of peritoneal macrophages to stimulation with TLR ligands and virus infection. Whereas the production of TNF and CXCL1 chemokine KC were comparable, macrophages from *LysM-Cre⁺;Akirin2^{fl/fl}* mice produced significantly less IL-6 and IL-12p40 than those from *LysM-Cre⁺;Akirin2^{fl/+}* mice upon stimulation with various TLR ligands (**Figure 1A**). In contrast, macrophages lacking Akirin1 did not show any defects in the production of cytokines to TLR stimulation (data not shown), indicating that Akirin2, but not Akirin1, plays a critical role in the regulation of proinflammatory cytokine production. It has been shown that Newcastle disease virus (NDV) and Encephalomyocarditis virus (EMCV) infection is recognized by RIG-I and MDA-5, respectively, in macrophages, resulting in the production of type I IFNs. *Akirin2*-deficient macrophages produced significantly less IL-6 and IFN- β than did WT cells after NDV and EMCV infection (**Figure 1B, Supplementary Figure S3**). Furthermore, Herpes simplex virus (HSV)-mediated production of IL-6 and IFN- β was also severely impaired in the absence of Akirin2 in macrophages. Since recognition of HSV is mediated through STING by cytoplasmic DNA sensors including cyclic GMP-AMP synthase (Sun et al, 2013), these results indicate that Akirin2 plays a pivotal role in the production of

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IL-6, IL-12p40 and IFN- β in response to the recognition of pathogens via various classes of sensors in macrophages.

Next we examined whether Akirin2 regulates IL-6 production at the level of gene expression. Northern blot analysis revealed that LPS-induced expression of genes encoding IL-6 and CXCL10 was severely impaired in LysM-Cre⁺;Akirin2^{fl/fl} macrophages relative to control LysM-Cre⁺;Akirin2^{fl/+} cells, indicating that Akirin2 is critical for the expression of several LPS-inducible genes (**Figure 1C**). However, the induction of genes encoding IFIT2 and CXCL1 chemokine *KC* was similar in LysM-Cre⁺;Akirin2^{fl/+} and LysM-Cre⁺;Akirin2^{fl/fl} macrophages. Gene induction in response to Poly I:C stimulation was similarly impaired in LysM-Cre⁺;Akirin2^{fl/fl} macrophages (**Figure 1D**). Time course analysis revealed that LPS- and Poly I:C-induced *Il6*, *Il12* and *Ifnb*, but not *Cxcl1* expression was altered at all time points tested in the absence of Akirin2 in peritoneal (**Supplementary Figure S4A, S4B**) and BM derived macrophages (BMDM) (**Supplementary Figure S5**). Thus, mouse Akirin2 regulates the expression of a set of LPS and Poly I:C inducible genes, including *Il6*, *Il12b* and *Ifnb*.

Akirin2 is critical for the responses to *Listeria monocytogenes* infection *in vivo*

Next we examined the innate immune responses to pathogen infection *in vivo* by using *Listeria monocytogenes* as a model. When LysM-Cre⁺;Akirin2^{fl/fl} mice were intraperitoneally infected with *Listeria monocytogenes*, the production of IL-6 and IL-12p40, but not TNF, in the sera was severely impaired compared with LysM-Cre⁺;Akirin2^{fl/+} mice (**Figure 2A**). Consistently, *Listeria*-mediated infiltration of myeloid cells including CD11b⁺Ly6G^{hi} neutrophils to the peritoneal cavity was significantly reduced in LysM-Cre⁺;Akirin2^{fl/fl} mice (**Figure 2B, 2C**). CD11b⁺Ly6c^{hi}Ly6g^{lo} inflammatory monocytes also tended to require Akirin2 for their recruitment (**Figure 2D**). Furthermore, the expression of *Il6*, but not *Tnf*, in the PECs from *Listeria*-infected mice decreased in the absence of Akirin2 in macrophages (**Figure 2E**). Consistent with the impaired immune responses to infection, Akirin2 deficiency leads to impaired clearance of infected *Listeria* in the spleen and liver (**Figure 2F**). These results

demonstrate that Akirin2 expressed in macrophages is critical for innate immune responses to *Listeria* infection and host defense *in vivo*.

Akirin2 is required for LPS-induced nucleosome remodeling at the *Il6* promoter region

As *Drosophila* Akirin acts together with or downstream of Relish, we next examined the LPS-dependent activation of NF- κ B in LysM-Cre⁺;Akirin2^{fl/fl} macrophages. In response to these stimuli, neither degradation of I κ B α (**Figure 3A**) nor induction of NF- κ B DNA binding (**Figure 3B**) was impaired in LysM-Cre⁺;Akirin2^{fl/fl} macrophages. In addition, LPS-induced phosphorylation of JNK and p38 was not altered between control and LysM-Cre⁺;Akirin2^{fl/fl} macrophages (**Figure 3A**). These data indicate that mouse Akirin2 acts together with or downstream of NF- κ B and MAP kinases in the control of TLR inducible gene expression in macrophages.

To examine if Akirin2 controls LPS-induced gene expression at the transcriptional level, we performed a chromatin-immunoprecipitation (ChIP) assay using anti-phospho-polymerase II (Pol II) Ab in LPS-stimulated macrophages. As shown in **Figure 3C**, phospho-Pol II was recruited to the *Il6* and *Cxcl1* promoters and coding regions in response to LPS stimulation in wild-type macrophages. However, its recruitment to the *Il6* locus was severely impaired in the absence of Akirin2 (**Figure 3C**). Furthermore, impaired recruitment of phospho-pol II to the *Il6* and *Il12b*, but not *Cxcl1*, promoter was observed at various time points after LPS stimulation (**Figure 3D**). Likewise, the recruitment of phospho-pol II to the *Tnf* promoter was normal in LysM-Cre⁺;Akirin2^{fl/fl} macrophages (**Supplementary Figure S6A**). Collectively, these results clearly indicate that Akirin2 controls *Il6* gene expression at the transcription level.

To comprehensively examine the effect of Akirin2 deficiency on LPS-induced gene expression in macrophages, we examined the genome wide change in gene expression in response to LPS by microarray analysis. We selected 1054 genes whose expression levels were upregulated more than 2-fold in response to LPS stimulation in wild-type macrophages (**Supplementary Table S1**). Comparison of LPS-inducible genes between wild-type and LysM-Cre⁺;Akirin2^{fl/fl} macrophages revealed that 187 genes were expressed less than 2-fold in

the absence of Akirin2. Hierarchical clustering of the 1054 genes divided the LPS-inducible genes into 8 different clusters based on their expression levels in LysM-Cre⁺;Akirin2^{fl/+} (Control) and LysM-Cre⁺;Akirin2^{fl/fl} (KO) macrophages (**Supplementary Table S1** and **Figure 3E**). Among these, gene clusters 1, 2 and 6 tended to be down-regulated in LysM-Cre⁺;Akirin2^{fl/fl} macrophages (**Figure 3E**). From these clusters we next selected 25 genes whose expression levels were highly different in wild-type and LysM-Cre⁺;Akirin2^{fl/fl} macrophages and designated these as Akirin2-dependent genes (**Figure 4A**). In addition, we selected 48 genes whose expression levels were comparable between wild-type and LysM-Cre⁺;Akirin2^{fl/fl} macrophages as a reference.

Using these two sets of genes, we first compared the promoter sequences of Akirin2-dependent and -independent genes. However, we did not observe any significant differences in the frequency of NF-κB binding sites between Akirin2-dependent and -independent gene promoters (**Supplementary Figure S7**). In contrast, the Akirin2-dependent genes tended to harbor significantly fewer CpG islands compared with Akirin2-independent genes ($p = 0.0067$), entire LPS-inducible genes ($p = 1.4 \times 10^{-6}$) or genes in the whole genome ($p = 0.0016$) (**Figure 4A, 4B**). These data demonstrate that among LPS-inducible genes Akirin2 controls the expression of a set of genes whose promoter regions have relatively fewer CpG islands, suggesting a relationship between Akirin2 and chromatin remodeling.

Akirin2 is critical for chromatin remodeling of the *Il6* promoter region

To investigate whether Akirin2 is involved in the regulation of chromatin remodeling, nuclei from unstimulated and LPS-stimulated peritoneal macrophages were treated with a restriction enzyme; after purification of genomic DNA and cleavage with a second enzyme, the efficiency of cleavage in the isolated nuclei was monitored by Southern blot (Zhou et al, 2004). Macrophages from LysM-Cre⁺;Akirin2^{fl/fl} mice showed decreased accessibility to the *Il6* promoter, indicating that Akirin2 is critical for opening the *Il6* promoter and further nucleosome remodeling (**Figure 4C**).

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We next investigated the involvement of Akirin2 in histone modification on the *Il6* promoter region in response to LPS. We examined changes in the trimethylation of histone 3 lysine 4 (H3K4) as well as acetylation of H3K9 on the *Il6*, *Il12b*, *Cxcl1* and *Tnf* promoters in response to LPS. Although trimethylation of H3K4 on the *Cxcl1* and *Tnf* promoters was comparably induced in response to LPS stimulation in LysM-Cre⁺;Akirin2^{fl/+} and LysM-Cre⁺;Akirin2^{fl/fl} macrophages, the *Il6* and *Il12b* promoters required Akirin2 for inducing H3K4 trimethylation (**Figure 4D, Supplementary Figure S6B**). Likewise, the acetylation of H3K9 on the *Il6* and *Il12b*, but not *Cxcl1* and *Tnf* promoters, was also induced in response to LPS in an Akirin2-dependent manner (**Figure 4E, Supplementary Figure 6C**). These results indicate that Akirin2 is directly involved in the regulation of chromatin remodeling and histone modification at the *Il6* promoter.

Akirin2 interacts with BAF60 family members, which are required for *Il6* expression

One mechanism whereby Akirin might function as a general cofactor for gene expression would be through interactions with chromatin remodeling complexes. A *Drosophila* whole-genome yeast 2-hybrid experiment (Giot et al, 2003) suggested that Akirin interacts with BAP60 (Brahma associated protein), a core subunit of the *Drosophila* SWI/SNF class Brahma (BRM) chromatin remodeling complex (Moller et al, 2005). Mammals have a family of BAP60 homologues, comprised of BAF60a, BAF60b and BAF60c, which are encoded by the *SMARCD1*, *SMARCD2* and *SMARCD3* genes, respectively (Puri & Mercola, 2012). Among these, BAF60c is preferentially expressed in the developing heart and somites during embryogenesis (Lickert et al, 2004). Furthermore, an equivalent activity of BAF60c in skeletal muscle differentiation and the molecular mechanism by which BAF60c promotes tissue-specific activation of gene expression has been described (Forcales et al, 2012). BAF60 proteins have been shown to interact with multiple transcription factors and are thought to bridge interactions between these transcription factors and SWI/SNF complexes, thereby allowing the recruitment of SWI/SNF to the target genes (Debril et al, 2004; Hsiao et al, 2003; Lickert et al, 2004).

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To investigate if Akirin2 interacts with mammalian BAF60 homologues, we overexpressed Flag-tagged Akirin2 and Myc-tagged BAF60 family members (BAF60a, BAF60b and BAF60c proteins) in HEK293 cells. Immunoprecipitation of the cell lysates with anti-Flag antibody co-precipitated Myc-tagged BAF60a, BAF60b and BAF60c (**Figure 5A**), indicating that Akirin2 can interact with BAF60 proteins in cells. Furthermore, endogenous Akirin2 co-precipitated with anti-Baf60a Ab in peritoneal macrophages irrespective of stimulation by LPS (**Figure 5B**), and the specificity of the Akirin2 precipitation was confirmed by the fact that the precipitated band was not observed in LysM-Cre⁺;Akirin2^{fl/fl} macrophages (**Figure 5C**). These results suggest that endogenous Akirin2 associates with Baf60 proteins even in resting cells.

To gain functional insight into the specific role of BAF60 proteins and their relation to Akirin2 in TLR/IL-1R-induced gene expression, we knocked down BAF60 genes, the SWI/SNF catalytic subunit BRG1 and AKIRIN2 in HeLa cells (**Supplementary Figure S8A**) and in J774 macrophages (**Supplementary Figure S8B and S8C**). The expression of *IL6* in response to IL-1 β was severely impaired in Akirin2, BAF60b, BAF60c and BRG1 depleted HeLa cells. In contrast, knockdown of BAF60a failed to affect *IL6* expression in response to IL-1 β stimulation (**Figure 5D**). However, the induction of genes encoding *IL8*, a counterpart of mouse *KC*, was comparable in control, Akirin2 and BAF60 depleted HeLa cells (**Figure 5E**). In addition, knockdown of Baf60 genes as well as Brg1 in the J774 mouse macrophage cell line led to the defects in the expression of *Il6*, but not *Cxcl1*, in response to LPS (**Supplementary Figure S8D**). In this cell line, Baf60a is also critical for the regulation of *Il6* in contrast to HeLa cells. These results indicate that Akirin2 and BAF60 complex members are critical for regulating *Il6* gene expression together with BRG1 in various cell types.

Although Akirin2 target genes have fewer CpG islands on their promoter regions, some of the target genes such as *Cmpk2*, *Akap12* and *Parp14* harbour a CpG island (**Figure 4A**). To investigate if these genes require chromatin remodelling for their expression, we examined their expression in Brg1 knockdown J774 macrophages in response to LPS. As shown in **Supplementary Figure S9**, LPS-induced expression of *Cmpk2*, *Akap12* and *Parp14* was

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impaired in Brg1 knockdown cells, indicating that Akirin2 target genes require SWI/SNF complex for their expression irrespective of the presence of CpG islands on their promoter regions.

Next we examined the role of BAF60 proteins in the regulation of histone modification. Knockdown of BAF60b and BAF60c, but not BAF60a, in HeLa cells resulted in deficient recruitment of phospho-Pol II as well as H3K9 acetylation in response to IL-1 β stimulation (**Figure 5F, 5G**). Similarly, downregulation of Akirin2 and Brg1 also impaired the recruitment of phospho-Pol II and acetylation of H3K9, further confirming that Akirin2, BAF60b and BAF60c regulate *Ilf6* gene expression through chromatin modification.

Finally, we investigated the interaction between Akirin2 and BAF60 family members at the residue level. The immunoprecipitation experiment further revealed that the Akirin2 mutants lacking N-terminal NLS (Δ 1-30) or the C-terminal region (Δ 151-203) were capable of co-precipitating with BAF60a or BAF60c (**Supplementary Figure S10A, S10B**), suggesting that Akirin2 interacts with BAF60 proteins via its central portion. Indeed, an alignment of 50 Akirin proteins among different species identified that the central portion is also highly conserved and is predicted to assume a helical secondary structure (**Figure 6A**).

Akirin2 interacts with I κ B- ζ via the C-terminal conserved region

We then further investigated the mechanism by which Akirin2 regulates gene expression in the course of TLR-induced macrophage activation. We hypothesized that Akirin2 acts in concert with NF- κ B family members by physically associating with them, though we did not observe interaction with NF- κ B family members (data not shown). On the other hand, it has been shown that *Ilf6* and *Ilf12b*, but not *Tnf*, mRNA expression to TLR stimulation requires an I κ B-like molecule called I κ B- ζ (Yamamoto et al, 2004; Yamazaki et al, 2001). Another report suggested the involvement of I κ B- ζ in the regulation of histone modifications, though the underlying mechanism is not well understood (Kayama et al, 2008). Interestingly, Flag-tagged I κ B- ζ coprecipitated with Myc-tagged Akirin2, when overexpressed in HeLa cells (**Figure 6B**). Furthermore, immunoprecipitation of overexpressed flag-tagged Akirin2 with anti-Flag Ab

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resulted in the co-precipitation of endogenous NF- κ B p50, but not p65, subunit in HeLa cells upon IL-1 β stimulation (**Figure 6C**). Interestingly, overexpressed Akirin2 could also precipitate endogenous I κ B- ζ with and without stimulation (**Figure 6C**). When Flag-Akirin2 was abundantly expressed, the interaction between Flag-Akirin2 and I κ B- ζ appears to be modestly increased probably due to the increase in the expression of I κ B- ζ . Consistently, even endogenous Akirin2 co-precipitated with endogenous I κ B- ζ in peritoneal macrophages (**Figure 6D**), and this interaction was not observed in I κ B- $\zeta^{-/-}$ or LysM-Cre⁺;Akirin2^{fl/fl} macrophages (**Figure 6D**). Whereas I κ B- ζ expression was induced in macrophages in response to LPS stimulation, the I κ B- ζ -Akirin2 interaction was observed even in resting cells and modestly decreased in response to LPS stimulation (**Figure 6D**). This may be due to the modest decrease in the endogenous Akirin2 protein expression in response to LPS stimulation in macrophages (**Supplementary Figure S1B**). These results suggest that endogenous I κ B- ζ expressed in resting cells can interact with endogenous Akirin2 irrespective of TLR stimulation. Then we examined the region of Akirin2 responsible for interaction with I κ B- ζ (**Figure 6A**). Antibody staining of the HeLa cells expressing Akirin2 mutants clearly showed a strict dependence on the NLS (Δ 1-30) (**Figure 6E**). Whereas a Myc-tagged Akirin2 mutant lacking the N-terminal NLS co-precipitated with Flag-tagged I κ B- ζ , deletion of the C-terminal conserved region of Akirin2 abrogated the interaction between I κ B- ζ and Akirin2 upon overexpression (**Figure 6F**). Furthermore, endogenous I κ B- ζ co-precipitated the Flag-tagged full-length and Δ 1-30 Akirin2 mutant, but not the mutant lacking the C-terminal region (**Figure 6G**). Next, we examined regions of I κ B- ζ responsible for interaction with Akirin2. I κ B- ζ is comprised of an N-terminal NLS, a central NF- κ B p50 subunit binding domain, and a C-terminal ankyrin repeat domain (Motoyama et al, 2005). Although deletion of I κ B- ζ N-terminal region (Δ 1-179) did not affect the interaction with Akirin2, the mutant I κ B- ζ lacking the C-terminal Ankyrin-repeat region (Δ 439–718) failed to co-precipitate with Akirin2 (**Figure 6H**). These data indicate that Akirin2 interacts with I κ B- ζ , and the Akirin2 C-terminal conserved region is critical for the interaction with the Ankyrin repeat domain of I κ B- ζ . Next we investigated the relationship between I κ B- ζ , Akirin2 and BAF60 family members. Interestingly, the immunoprecipitation experiment

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revealed that overexpressed Baf60a co-precipitated with endogenous I κ B- ζ and this interaction is probably mediated through the endogenous Akirin2, as endogenous I κ B- ζ failed to co-precipitate Baf60a upon overexpression of the C-terminal deletion mutant of Akirin2 lacking the I κ B- ζ binding domain (**Figure 6I**). Collectively, these results demonstrate that I κ B- ζ can interact with Baf60 proteins via bridging by Akirin2.

The I κ B- ζ -Akirin2 cascade is critical for the recruitment of Brg1 to the *Il6* promoter for transactivation

Next we examined the role of the interaction between Akirin2 and I κ B- ζ . The C-terminal part of Akirin2 is highly conserved among species, although the functional role of this domain is enigmatic. Thus we investigated the role of the Akirin2 C-terminal region in response to TLR/IL-1R stimulation. Whereas deletion of Akirin2 in MEFs resulted in severe reduction of LPS-mediated expression of *Il6*, but not *Cxcl1*, consistent with a previous report (Goto et al, 2008), the expression of full length Akirin2 rescued *Il6* expression upon LPS stimulation at the mRNA and protein levels (**Figure 7A**). In contrast, the deletion of either the N-terminal or C-terminal resulted in defective rescue of IL-6 production in response to LPS or IL-1 β stimulation (**Figure 7B, 7C**). Thus, these results suggest that the interaction between I κ B- ζ and Akirin2 is critical for inducing a set of genes in response to TLR stimulation.

Since I κ B- ζ has been shown to be critical for the histone modification of its target genes in response to LPS, we next sought to determine if Akirin2 and I κ B- ζ are required for the recruitment of the SWI/SNF remodeling complexes to the promoter regions of their target genes. ChIP experiments were performed using anti-BRG1 antibody and sheared cross-linked chromatin was prepared from unstimulated or LPS-stimulated macrophage cells lacking Akirin2 or I κ B- ζ . BRG1 recruitment to the *Il6* and *Il12b*, but not *Cxcl1* promoter was observed 1, 4 and 8 hours after LPS stimulation, and was impaired in the macrophages from LysM-Cre⁺;Akirin2^{fl/fl} mice as well as from I κ B- ζ -deficient mice (**Figure 7D**). Thus, these results indicate that both Akirin2 and I κ B- ζ are required for the recruitment of the SWI/SNF complex to the *Il6* and *Il12b* promoter regions.

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Next we examined if Akirin2 is also recruited to the promoter regions of its target genes. We retrovirally expressed Flag-tagged Akirin2 in wild-type and *NF-κBp50*^{-/-} BM macrophages, and performed the ChIP assay using anti-Flag Ab upon LPS stimulation (**Supplementary Figure S11A, S11B**). The levels of total Akirin2 mRNA expression increased modestly in Flag-Akirin2 expressing cells (**Supplementary Figure S11A**). Interestingly, we observed the recruitment of Akirin2 to the *Il6* and *Il12b*, but not *Cxcl1*, promoters in response to LPS stimulation (**Figure 7E**). In contrast, retroviral expression of Flag-tagged Akirin2 in *IκB-ζ*^{-/-} and *NF-κBp50*^{-/-} BM macrophages led to the abrogation of the recruitment of Akirin2 on the *Il6* and *Il12b* promoters (**Figure 7E and 7F**), indicating that IκB-ζ is required for the recruitment of Akirin2 to its target gene promoters. We further examined the recruitment of IκB-ζ to the immune regulatory gene promoters by retrovirally expressing Flag-tagged IκB-ζ in LysM-Cre⁺; *Akirin2*^{fl/fl} BM macrophages (**Supplementary Figure S11C**). The ChIP assay revealed that IκB-ζ is also recruited to the *Il6* and *Il12b*, but not *Cxcl1*, promoters in response to LPS stimulation (**Figure 7G**). Interestingly, the recruitment of IκB-ζ to the promoters required the presence of Akirin2 in macrophages (**Figure 7G**), suggesting that IκB-ζ and Akirin2 function together in order to be recruited to the *Il6* and *Il12b* promoters for inducing the gene expression.

Discussion

In *Drosophila*, Akirin is critical for gene expression downstream of the Imd pathway. We have previously shown that Akirin2, but not Akirin1, controls IL-6 production in mouse embryonic fibroblasts, probably downstream of NF- κ B activation. However, the early embryonic lethality of Akirin2-deficient mice prevented examination of the role of this molecule in innate immune cells. Furthermore, the mechanisms of Akirin2-mediated control of IL-6 production have not been clarified. In the present study, we investigated the role of Akirin2 in macrophages by generating an *Akirin2* conditional allele in mice, and we found that this molecule plays an essential role in the regulation of inflammatory gene expression in response to TLR stimulation as well as infection with viruses and a model bacteria, *Listeria monocytogenes*. Furthermore, Akirin2 in macrophages and neutrophils plays a role in the clearance of infecting bacteria. Thus, Akirin2 expressed in macrophages and neutrophils is critical for host defense against bacterial infection.

Akirin2 controls the TLR-induced transcription of a set of inflammation-related genes such as *Il6*, *Il12b* and *Ifnb*. In contrast, *Tnf* and *Cxcl1* are not regulated by Akirin2. It has been shown that *Tnf* gene expression is rapidly inducible compared with *Il6* or *Il12b*. *Cxcl1* was also reported to be a rapidly inducible gene in response to TNF stimulation. Therefore, it is suggested that Akirin2 is critical for the control of rather slowly inducible genes in the course of macrophage activation.

Promoters of Akirin2-dependent genes harbor significantly lower frequencies of CpG islands compared with Akirin2-independent genes, although there are some exceptions. Akirin2 is essential for the recruitment of the chromatin remodeling complex component Brg1 and histone modification in response to LPS stimulation. Nevertheless, CpG-containing Akirin2-dependent genes also require Brg1 for their expression, further supporting a role for Akirin2 in the control of genes that require chromatin remodeling for their expression. Mechanistically, we found that Akirin2 interacts with BAF60 proteins and I κ B- ζ , and the complex is recruited to *Il6* and *Il12b* promoters for inducing chromatin remodeling. Akirin2

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and I κ B- ζ are recruited to the *Il6* and *Il12b* promoters, and act together to recruit the SWI/SNF complex to the promoters of their target genes.

Dynamic changes in gene transcription in innate immune cells is pivotal for evoking inflammation in response to pathogen infection. Gene expression is, in turn, controlled by elaborate and complex mechanisms. Whereas the activation of transcription factors such as NF- κ B, AP-1 and IRF-3/-7 is essential for the induction of gene expression, the kinetics and intensity of the change in gene transcription is also regulated by histone modification and changes in chromatin architecture. TLR-inducible genes have been roughly classified into rapidly induced genes and late induced genes. The rapidly induced genes are known to be induced by the action of NF- κ B, AP-1 and IRF transcription factors, and frequently have promoters containing CpG islands. TLR signaling leads to rapid recruitment of NF- κ B as well as basic transcriptional machinery, p-TEFb phosphorylating Pol II to start generation of mature transcripts. On the other hand, the late induced genes are reported to be comprised of both CpG and low CpG island promoters (Bhatt et al, 2012). Low CpG island promoters are enriched in the highly induced genes, and possibly require chromatin remodeling for transcription factor binding and general transcription initiation. Interestingly, genes regulated by the presence of Akirin2 show quite low numbers of CpG islands and chromatin remodeling was impaired in the absence of Akirin2. We further found that Akirin2 is critical for recruiting the SWI/SNF complex to secondary gene promoters. These data demonstrate that Akirin2 functions as an adaptor molecule to tether the chromatin remodeling machinery to the inflammation related gene promoters. Akirin is highly conserved among orthologs identified throughout metazoa including *Drosophila*, mice, teleosts, and humans (Macqueen & Johnston, 2009). The function of the Akirin protein has remained enigmatic; despite its high degree of conservation, it remains almost completely devoid of known functional domains, catalytic activity, or demonstrable DNA binding (Goto et al, 2008; Macqueen & Johnston, 2009; Nowak et al, 2012). Akirin2 contains two conserved helical regions, one located in the central region (residues 75-95) and one in the C-terminus (residues 143-193), separated from each other and from the N-terminus by regions predicted to be intrinsically disordered. Neither of the conserved helical regions

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show significant sequence similarity to proteins of known function, but conserved intrinsically disordered regions are often important for mediating protein-protein interactions (Dyson & Wright, 2005). Akirins function during immune and inflammatory responses in *Drosophila* as well as in mice. Importantly, Akirins appear to have conserved functions in both the *Drosophila* and mouse immunity pathways, reinforcing the notion that the role of the Akirin protein has been conserved throughout evolution. Our results give new insight into the role of Akirin2 in mediating NF- κ B dependent gene expression in a SWI-SNF dependent manner via I κ B- ζ during inflammatory responses.

Akirin2 can interact with all 3 Baf60 proteins. Knockdown of all of the Baf60 proteins led to impaired *Il6* gene expression in response to LPS in the J774 macrophage cell line, though BAF60b and BAF60c, but not BAF60a, were important for *Il6* gene expression in response to IL-1 β in HeLa cells. Although the function of BAF60a has not been well understood, downregulation of BAF60b and c have been reported to reduce the expression of certain muscle genes, such as muscle creatine kinase (MCK) (Forcales et al, 2012) and this peculiarity has been ascribed to the partial, functional redundancy between BAF60b and c (Takeuchi & Bruneau, 2009). Based on this evidence, as well as on our initial identification by co-immunoprecipitation experiments that Akirin2 interacts with both BAF60b and c, we restricted our analysis to HeLa cells in which Akirin2, Brg1, BAF60b or c were individually knocked down. BAF60b and BAF60c have been shown to be modified while regulating chromatin remodeling. Besides muscle cell development, BAF60c was reported to be involved in lipogenesis (Wang et al, 2013). Insulin signaling leads to the phosphorylation of BAF60c by atypical protein kinase C ζ/λ inducing its nuclear translocation and regulation of lipogenic genes. On the other hand, BAF60b was reported to interact with an E3 ubiquitin ligase Unkempt, and ubiquitination of BAF60b results in its degradation in a proteasome-dependent fashion (Lores et al, 2010). In this regard, it is also interesting to explore if the BAF60 proteins undergo modification in the course of inflammation via Akirin2. Further studies are required to address differential involvement of the BAF60 proteins in controlling inflammation *in vivo*.

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In this study, we found that I κ B- ζ was required for the recruitment of Akirin2 as well as Brg1 to the *Il6* and *Il12b* promoters in macrophages in response to LPS. Reciprocally, Akirin2 is also required for the recruitment of I κ B- ζ to the same promoters in response to LPS. The interaction between endogenous I κ B- ζ and Akirin2 was also observed in resting macrophages. In addition, I κ B- ζ was found to form a complex with Baf60 proteins via Akirin2. These data suggest that I κ B- ζ is recruited to the LPS-inducible gene promoters together with Akirin2 and BAF60 proteins, where they induce chromatin remodeling. Since I κ B- ζ has been shown to interact with NF- κ B p50 subunit (Motoyama et al, 2005; Yamamoto et al, 2004), and NF- κ B p50 translocate from the cytoplasm to the nucleus upon IL-1 β /TLR stimulation, our results indicate that the complex of I κ B- ζ -Akirin2-SWI/SNF interacts with the NF- κ B p50 subunit via I κ B- ζ upon IL-1 β /TLR stimulation and the whole complex is then recruited to the *Il6* and *Il12b* promoters. Therefore, it is possible that the specificity of Akirin2-dependent and -independent genes is determined by the recruitment of the NF- κ B p50 subunit to the promoters of its target genes. Future studies will identify the recruitment of NF- κ B p50 subunit, I κ B- ζ and Akirin2 to each gene promoter in a comprehensive manner.

We have previously shown that I κ B- ζ functions in the production of IFN- γ in natural killer (NK) cells in response to IL-12 and IL-18 stimulation. NK cells required I κ B- ζ for inducing histone modifications in response to IL-12 and IL-18 stimulation (Miyake et al, 2010), suggesting that I κ B- ζ also controls chromatin remodeling in NK cells. It has also been shown that I κ B- ζ controls Th17 differentiation in T cells by regulating IL-17 gene expression via its promoter (Okamoto et al, 2010). In contrast, I κ B- ζ expressed in epithelial cells of lacrimal glands is critical for the prevention of their apoptotic cell death, which leads to a pathology similar to Sjogren's syndrome (Okuma et al, 2013), though its molecular mechanism is yet to be clarified. It is intriguing to speculate that I κ B- ζ functions via Akirin2-mediated chromatin remodeling in various immune cell types. Further studies with conditional Akirin2-deficient mice will uncover the broad functional roles of Akirin2 in various cells types *in vivo*.

In addition to the regulation of *Il6* and *Il12b* in response to stimulation with TLR ligands, Akirin2 was found to be important for the induction of type I IFNs and a set of IFN-inducible

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genes in response to RNA virus infection. It has been shown that a nucleosome masks the TATA box and the transcription start site of the human IFNB promoter, and that the nucleosome slides due to the action of SWI/SNF complex, enabling transcription (Agalioti et al, 2000). Further, a set of IFN-inducible genes require chromatin remodeling by SWI/SNF, a complex that includes a specific subunit, BAF200 (Yan et al, 2005). Thus, the involvement of Akirin2 in the regulation of IFN-inducible genes such as IL-6 in response to RLR stimulation can be explained by the regulation of chromatin remodeling of their promoter regions.

In addition to the control of innate immune responses, *Drosophila* Akirin has been shown to be required for Twist-mediated gene transcription by controlling chromatin remodeling during embryogenesis. Akirin mutant embryos had muscle defects in fly. Given that the lack of Akirin2 in mice results in early embryonic lethality, it will be interesting to investigate the role of Akirin2 in developmental tissues for controlling chromatin remodeling by coupling with various transcription factors.

In summary, this study clearly demonstrates that Akirin2 controls a set of inflammatory genes with non-CpG island promoters by recruiting the SWI/SNF complex and interacting with I κ B- ζ . We believe that this is the first study unveiling the mechanism of initiation of chromatin remodeling after NF- κ B activation in innate immune cells. Targeting Akirin2 in immune cells might be beneficial for ameliorating inflammatory diseases and may lead to new strategies for combating autoimmune diseases.

Experimental Procedures

Mice, Cells and reagents

Floxed *Akirin2*, *IκBζ*^{-/-} and *NF-κBp50*^{-/-} mice were described previously (Goto et al, 2008; Sha et al, 1995; Yamamoto et al, 2004). Mice were housed in specific-pathogen-free conditions and all animal experiments were done with the approval of the Animal Research Committee of the Research Institute for Microbial Diseases (Osaka University, Osaka, Japan), and of Kyoto University.

Peritoneal exudate cells were isolated from the peritoneal cavities of mice 3 days after injection with 2 ml of 4.0% Brewer's thioglycollate medium (Sigma) by washing with ice-cold Hank's buffered salt solution (Invitrogen). BMDMs were generated in RPMI-1640 medium containing 10% (vol/vol) FCS, 50 μM 2-mercaptoethanol and 20 ng/ml M-CSF (Peprotech). J774 mouse macrophage cell line was cultured and maintained in DMEM containing 10% (vol/vol) FCS, 50 μM 2-mercaptoethanol, 100 U/ml penicillin and 100 μg/ml streptomycin. TLR ligands, MALP-2 was provided as described previously (Matsushita et al, 2009), LPS from *Salmonella Minnesota* Re595 (Sigma), poly (I:C) from Amersham Biosciences; and R-848 and CpG oligonucleotide (ODN 1668) was used as described previously (Matsushita et al, 2009). Lipofectamine 2000, Lipofectamine RNAiMAX Reagent (Invitrogen) and FuGENE Transfection reagent (Promega) were used for transfection. Recombinant human IL-1β, mouse IL-6, IL-3 and stem cell factor was purchased from R&D systems.

Measurement of cytokine production

Peritoneal macrophages (5×10^4) and BMDM (5×10^4) were stimulated for 24 hours with Malp2, R848, LPS, Poly I:C or with CpG DNA. Culture supernatants were collected and cytokine concentrations were measured with an ELISA kit (R&D Systems) for IL-6, TNF, IL-12p40, CXCL1 and (PBL interferon source) for IFN-β.

In vivo infection experiments with *Listeria monocytogenes*

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L. monocytogenes bacteria were grown in Tryptic Soy Broth (TSB) at 37°C for 6 hours with shaking. Mice were intraperitoneally infected with 5×10^5 bacteria/mice in a volume of 0.25 ml of PBS. Serum was collected at the indicated time intervals and subsequently analyzed for cytokine concentrations with an ELISA kit (R&D Systems) for IL-6, TNF and IL-12p40 production. Mice were sacrificed 24 hours after infection, PECs were harvested and subjected to quantitative PCR and flow cytometry analysis. For measuring the bacterial load, liver and spleen were aseptically removed from mice to produce a homogenized PBS suspension. The serially 10-fold diluted homogenates by PBS of organs were placed on TSB agar plates and then were incubated at 37°C for 24 h. The number of bacterial colonies was then counted and expressed as CFU/organ.

Electrophoretic mobility-shift assay (EMSA)

Macrophage from the peritoneal exudate cells (1×10^6) were treated with LPS for various time periods. Nuclear extracts were purified from cells, incubated with probes specific for the NF- κ B and IRES DNA-binding sites, separated by electrophoresis and were visualized by autoradiography as described previously (Goto et al, 2008).

Co-Immunoprecipitation

HEK293 or HeLa cells seeded on 60- or 100-mm dishes were transiently transfected with a total of 6–8 μ g of the appropriate combination of plasmids. At 48 hours after transfection, cells were lysed in lysis buffer (20 mM Tris-HCl, pH 7.5, 150 mM NaCl, 1 mM EDTA and 1% (vol/vol) Nonidet P-40) containing complete mini protease inhibitor cocktail (Roche). Proteins were immunoprecipitated from lysates incubated overnight with an anti-flag-M2 monoclonal antibody (Sigma) and dynabeads (Invitrogen) in lysis buffer. Immune complexes were washed three times with lysis buffer and suspended in SDS sample buffer (lysis buffer containing 3 mM Tris-HCl, pH 6.8, 2% (wt/vol) SDS, 5% (vol/vol) β -mercaptoethanol, 10% (vol/vol) glycerol and bromophenol blue). Samples were boiled for 5 min at 98 °C and separated by

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SDS-PAGE and analyzed by immunoblot using anti-flag, anti-myc (Sigma), anti-NFκB-p50 (Santa Cruz H-119), anti-NFκB-p65 (Santa Cruz C-20) or anti-IκB-ζ antibodies.

Immunoblot analysis

Peritoneal exudate cells were cultured for 2 hour in medium and were then stimulated with LPS for the indicated times and were lysed with lysis buffer (20 mM Tris-HCl, pH 7.5, 150 mM NaCl, 1 mM EDTA and 1% (vol/vol) Nonidet P-40) containing complete mini protease inhibitor cocktail (Roche). Cell lysates were separated by standard SDS-PAGE and analyzed by immunoblot. Antibodies to the following proteins were used: phosphorylated Erk (Cell Signaling no. 9101), phosphorylated p38 (Cell Signaling 9211), p38 (Santa Cruz C-20), Erk (Santa Cruz K-23), IκBα (Santa Cruz C-21), phosphorylated JNK (Cell Signaling 9251), JNK (Santa Cruz C-17) and β-tubulin (Santa Cruz D-10).

Quantitative PCR analysis

Total RNA was isolated with TRIzol (Invitrogen), and reverse transcription was performed with ReverTra Ace (Toyobo) according to the manufacturer's instructions. For quantitative PCR, cDNA fragments were amplified by real time PCR Master Mix (Toyobo); fluorescence from the TaqMan probe for each cytokine was detected by a 7500 real-time PCR system (Applied Biosystems). To determine the relative induction of cytokine mRNA in response to various stimuli, the mRNA expression level of each gene was normalized to the expression level of 18S rRNA. All the experiments were performed in triplicate at least three times.

Flow cytometry

Antibodies for flow cytometry were purchased from BD Biosciences and Biolegend. Cells were washed in ice-cold flow-cytometry buffer (0.5% (vol/vol) FCS and 2 mM EDTA in PBS, pH 7.5), then incubated with each antibody for 30 min, washed twice with flow-cytometry buffer and resuspended in an appropriate volume of flow-cytometry buffer. Data were acquired on a FACS Canto II flow cytometer (BD Biosciences) and analyzed with FlowJo (Tree Star).

Restriction enzyme accessibility assay

Experiments were performed as described previously with little modifications (Ramirez-Carrozzi et al, 2006). Briefly, PECs (2×10^6 cells) were stimulated with LPS (100 ng/mL) for indicated time period and the cell nuclei were isolated and digested with limiting amounts of restriction enzyme *Afl*III (100 U), and were incubated for 15 min at 37°C, followed by genomic DNA isolation. Purified DNA (10–15 μ g) was digested to completion to generate reference cleavage products using the following restriction enzymes: *Xba*I and *Spe*I for the *Il6* promoter. Samples were analyzed by Southern blotting with 32 P-labeled gene-specific probes designed at the *Il6* promoter region (-1169 to +620).

Chromatin Immunoprecipitation (ChIP)

ChIP was performed by using a chip assay kit (Millipore) as described elsewhere with little modification (Sato et al, 2010). Briefly, PECs or BMDM's (2×10^6 cells) and HeLa cells (5×10^6) were stimulated with LPS (1 μ g/mL) or recombinant hIL-1 β (10 ng/ml) (R&D systems), respectively, and were fixed with 1% formaldehyde (Nacalai Tesque, Japan) for 10 min at 37°C. Cells were then washed twice with ice cold PBS and resuspended in SDS lysis buffer supplied with the kit. Lysates were sonicated using ultrasonicator (Covaris S2 and Bioruptor™) to obtain DNA fragments with a peak in size between 150 and 300 bp. Lysates pre-cleared with Dynabeads (Invitrogen) were incubated with antibodies against phospho RNA Pol II (S5P), H3K9(Ac) or H3K4(Me)3 (Abcam) and Brg-1 (J1) and immune-precipitated at 4°C overnight. The immune complexes were absorbed with beads. Beads were washed once with low salt buffer, high salt buffer, LiCl wash buffer, and twice with TE buffer. Immune complexes were extracted with elution buffer (50mM Tris-HCl, pH-8.0; 10mM EDTA; 1% SDS) and cross links were reversed by incubating overnight at 65°C. After RNase treatment for 2 hours at 37°C followed by proteinase K treatment for 2 hours at 56°C, DNA was then purified via phenol-chloroform precipitation. The purified DNA was quantified and then used for qPCR analysis

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to assess the presence of target sequences. Quantitative RT-PCR was performed with SYBR green qPCR mix (Toyobo) in an Applied Biosystem 7500. Primers used for amplifying *Il6*, *KC* (*Cxcl1*), *Tnf*, *Il12b* and *IL8* loci were shown previously (Coles et al, 2010; Negishi et al, 2012; Ramirez-Carrozzi et al, 2006). ChIP values were normalized against the input and expressed as relative enrichment of the material precipitated by the indicated antibody on *Il6* promoter (relative quantification using the comparative Ct method ($2^{-\Delta\Delta C_t}$)). Error bars indicate mean \pm std. deviation. The results are representative of at least three independent experiments.

Construction of expression plasmids

Akirin2, I κ B ζ and Baf60a, b and c cDNA was obtained by PCR from a mouse cDNA library. Full length or truncated Akirin2 cDNAs were cloned into the pcDNA-Flag or Myc tag vectors for overexpression or pMRX- Puro vector for retrovirus production (Matsushita et al, 2009). Full length Baf60a, b and c cDNAs were cloned into pcDNA3.1 (+) -Myc tag vectors and full length or truncated I κ B ζ cDNAs were cloned into the Flag tagged pEF-BOS vector for overexpression or pMRX- Puro vector for retrovirus production.

Retroviral transduction of Mouse Embryonic Fibroblasts (MEFs)

Akirin2^{-/-} MEFs were established as described (Goto et al, 2008). Briefly, MEFs were isolated from *Akirin2*^{fl/fl} mice at embryonic day 13.5. Cells were cultured in DMEM (supplemented with 10% (vol/vol) FCS, 10 mM sodium pyruvate, 2 μ M L-glutamine, 50 μ M 2-mercaptoethanol, 100 U/ml penicillin, 100 μ g/ml streptomycin). Then, 48 h later, these cells were transduced with retroviral supernatant (supplemented with 10 ng/ml of polybrene) for 12 hours. Virus was produced by PlatE packaging cells transfected with pMRX-puro vector stably expressing -cre- and/or full length Akirin2 or deletion mutants. After incubation with retroviral supernatant (supplemented with polybrene), the cells were cultured in DMEM (supplemented with 10% (vol/vol) FCS, 10 mM sodium pyruvate, 2 μ M L-glutamine, 50 μ M 2-mercaptoethanol, 100 U/ml penicillin, 100 μ g/ml streptomycin and 2 μ g/ml of puromycin).

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After 48 hours in culture, cells were washed once with ice cold PBS and the cells were harvested and used for further experiments.

Retroviral transduction of Macrophages

Highly efficient retroviral transduction of macrophages was achieved by transduction of hematopoietic stem cells before differentiation into macrophages, as described previously (Sato et al, 2010). BM was isolated from $\text{LysM-Cre}^+; \text{Akirin2}^{\text{fl/+}}$, $\text{LysM-Cre}^+; \text{Akirin2}^{\text{fl/fl}}$, $\text{NF-}\kappa\text{Bp50}^{-/-}$ and $\text{IkB}\zeta^{-/-}$ deficient mice that had been injected intraperitoneally 4 d earlier with 5 mg of 5-fluorouracil (Nacalai tesque). Cells were cultured in stem cell media (DMEM supplemented with 15% FCS, 10 mM sodium pyruvate, 2 mM L-glutamine, 50 μM β -mercaptoethanol, 100 U/ml penicillin, 100 $\mu\text{g/ml}$ streptomycin, 100 ng/ml stem cell factor, 10 ng/ml IL-6 and 10 ng/ml IL-3). Then, 48 h later, these cells were transduced with retroviral supernatant (supplemented with stem cell factor, IL-6, IL-3 and 10 ng/ml of polybrene) on two successive days. Virus was produced using PlatE. After the second transduction, cells were washed and resuspended in macrophage growth media (RPMI 1640 medium supplemented with 10% FCS, 10 mM HEPES, 10 mM sodium pyruvate, 2 mM L-glutamine, 50 μM β -mercaptoethanol, 100 U/ml penicillin, 100 $\mu\text{g/ml}$ streptomycin and 40 ng/ml macrophage colony-stimulating factor). The cells were cultivated for 7 days, and were subjected to analysis.

Microarray Analysis

Thioglycollate-elicited peritoneal macrophages were stimulated with LPS (100 ng/mL) for different time periods. Then, total RNA was extracted with TRIzol (Invitrogen Life Technologies) and further purified using an RNeasy kit (Qiagen). Biotin-labeled cDNA was synthesized from 100 ng of total RNA using the Ovation Biotin RNA amplification and labeling system (Nugen) according to the manufacturer's protocol. Hybridization, staining, washing, and scanning of Affymetrix mouse Genome 430 2.0 microarray chips were conducted according to the manufacturer's instructions. Data analysis was performed using R (<http://www.r-project.org/>). The data were normalized by robust multichip analysis (RMA) and

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log2-transformed expression values are shown. Genes with differential expression between control and *Akirin2*-deficient macrophages were clustered based on the similarity in the differences in microarray probe intensities between control and *Akirin2*-deficient macrophages for the time points 0, 2, and 4 h after stimulation. Clustering was done using hierarchical clustering using Ward's method. After visual inspection genes were divided into 8 clusters. MicroArray data are deposited in GEO (Accession number GSE59319).

Analysis of CpG islands associated with *Akirin2*-dependent and independent genes

For *Akirin2*-dependent and independent genes, the GC content and CpG score of the genomic regions -100 to -1 and +1 to +100 relative to the transcription start sites were calculated. CpG island predictions for the mouse genome were obtained from the UCSC Genome Browser database (Meyer et al, 2013). For the *Akirin2*-dependent and independent sets of genes, the number of genes with predicted CpG islands in the regions -1kb to +1kb and -10kb to +10kb relative to the transcription start site were counted. P values for the difference in CpG island-associations between the *Akirin2*-dependent and independent sets were estimated using the Z-test for proportions, based on the predicted CpG-islands in the region -1kb to +1kb. All analyses were based on the Refseq gene annotations for version mm10 (Dec. 2011, GRCm38) of the mouse genome.

RNA Interference

siRNAs targeting human AKIRIN2, BAF60a, BAF60b and BAF60c were designed and purchased from the Ambion (**Supplementary Table S2**). siRNAs targeting human BRG1 were designed and purchased from Dharmacon (**Supplementary Table S2**). HeLa cells were transfected with siRNA using Lipofectamine RNAiMax following the manufacturer's instructions with little modifications. A retroviral vector containing a small hairpin RNA (shBAF60a, shBaf60b, shBaf60c and shBrg1) was obtained by cloning the randomly designed annealed primers into BglII and XhoI sites of the pSUPER-retro-Puro (Oligoengine) following

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the manufacturer's protocol. J774 cells were transfected with shRNA using FuGENE transfection reagent according to the manufacturer's instructions with little modifications.

Statistical Analysis

The statistical significance of differences was determined by the two-tailed Student's *t*-test. Differences with a *P* value of less than 0.05 were considered statistically significant.

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Acknowledgement

We thank all the colleagues in our laboratory for helpful discussion and suggestions. E. Kamada, M. Kageyama and Y. Asahira for secretarial assistance. Y. Fujiwara and M. Kumagai, for technical assistance. We thank Dr. Tatsushi Muta (Tohoku University) and Dr. Weidong Wang (NIH) for kindly providing anti-I κ B- ζ and anti-Brg1 (J1) antibody, respectively. We also thank Dr. Masao Mitsuyama (Kyoto University) for kindly providing *Listeria monocytogenes*. *NF- κ Bp50^{-/-}* mice were a kind gift from Dr. Eiijiro Jimi (Kyushu Dental University). This work was supported in part by the Japan Society for the Promotion of Science (JSPS) through the Funding Program for World-Leading Innovative R&D on Science and Technology (FIRST Program), and by JSPS Core-to-Core Program, A. Advanced Research Networks. This work was also supported in part by grants from Takeda Science Foundation, Naito Foundation, and Mochida Foundation. S.T. is supported by a research fellowship from the MEXT for the Promotion of Science for Young Scientists. J.-M.R. was supported by Centre National de la Recherche Scientifique, grants from Agence Nationale de la Recherche (LABEX : ANR-10-LABX-0036_NETRNA) and the European Research Council (Immunodroso2009-AdG-20090506).

Author Contributions

S.T. performed most of the experiments. S.T. and O.T. designed experiments, analyzed the data and wrote the manuscript. A.V. performed bioinformatics analysis. D.M.S. performed structural modeling. K.M. helped with the experiments. D.O., T.I. and T.M. participated in discussions. J.M.R. and J.A.H. shared information on *Drosophila* and mammalian Akirin. S.A. and O.T. supervised the project.

Competing Interests Statement

The authors declare that they have no competing financial interests.

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Figure Legends

Figure 1: TLR-ligand- and virus infection-induced cytokine production and gene expression in *Akirin2*-deficient macrophages.

Peritoneal macrophages from *LysM-Cre⁺;Akirin2^{fl/+}* and *LysM-Cre⁺;Akirin2^{fl/fl}* mice were stimulated with various TLR ligands including Malp2 (1, 10 ng/ml), poly I:C (10, 100 µg/ml), LPS (10, 100 ng/ml), R848 (10 ng/ml) and CpG-DNA (0.1, 1 µM) (A) or infected with HSV and NDV (MOI 1, 5) (B) for 24 h. Then, IL-6, TNF, CXCL-1, IL-12p40 and IFN-β concentrations in the culture supernatants were determined by ELISA. Peritoneal macrophages from *LysM-Cre⁺;Akirin2^{fl/+}* and *LysM-Cre⁺;Akirin2^{fl/fl}* mice were stimulated with 100 ng/ml LPS (C) or poly I:C (200 µg/ml) (D) for indicated time period and the total RNA extracted was subjected to Northern blot analysis for the expression of *Il6*, *Ifnb*, *Cxcl1*, *Ifi2*, *Cxcl10* and *Actb*. Error bars indicate mean ± std. deviation. Results are representative of at least three independent experiments. Statistical significance was determined using the Students *t*-test. **p* < 0.05; ***p* < 0.01.

Figure 2: *Akirin2* is critical for innate immune responses to *Listeria* infection *in vivo*

LysM-Cre⁺;Akirin2^{fl/+} and *LysM-Cre⁺;Akirin2^{fl/fl}* mice (n=7) were intraperitoneally infected with 5×10^5 *L. monocytogenes* /mice in a volume of 0.25 ml of PBS. (A) Serum cytokine levels were determined by ELISA for IL-6, TNF and IL-12p40 at indicated time periods following infection. (B) PECs were prepared from *LysM-Cre⁺;Akirin2^{fl/+}* (n=4) and *LysM-Cre⁺;Akirin2^{fl/fl}* (n=5) mice 24 h after *Listeria* infection, and the cell surface expression of myeloid cell markers was determined by flow cytometry. Frequencies of CD11b⁺Ly6g^{hi} neutrophils and CD11b⁺Ly6c^{hi}Ly6g^{lo} inflammatory monocytes are indicated. (C) The numbers of total PECs, CD11b⁺ myeloid cells and (D) CD11b⁺Ly6c^{hi}Ly6g^{lo} inflammatory monocytes and CD11b⁺Ly6g^{hi} neutrophils are shown. (E) The levels of *Il6* and *Tnf* mRNAs in the PECs were measured by Q-PCR. (F) The bacterial load in the liver and spleen were measured by colony formation assay and the number of bacterial colonies was then counted and expressed

as CFU/organ (n=4). Error bars indicate mean \pm std. deviation. Statistical significance was determined using the Students *t*-test. **p* < 0.05; ***p* < 0.01; ****p* < 0.001.

Figure 3: Akirin2 is dispensable for NF- κ B activation but decisive for the transcription progression

(A) Whole cell lysates were prepared from LysM-Cre⁺;Akirin2^{fl/+} and LysM-Cre⁺;Akirin2^{fl/fl} PECs treated with LPS (100 ng/ml) for the indicated time period and I κ B α , β -tubulin, p-JNK, JNK, p-p38, p38, p-ERK and ERK expression were determined by western blotting. (B) Nuclear extracts were prepared from LysM-Cre⁺;Akirin2^{fl/+} and LysM-Cre⁺;Akirin2^{fl/fl} PECs treated with LPS (100 ng/ml) for the indicated time period and NF- κ B- and ISRE-DNA binding activity were analyzed by Electrophoretic Mobility-Shift Assay (EMSA). (c, d) ChIP experiments were performed with chromatin prepared from LysM-Cre⁺;Akirin2^{fl/+} and LysM-Cre⁺;Akirin2^{fl/fl} PECs treated with or without LPS (1 μ g/ml) for indicated time period. Antibody against phospho-RNA polymerase II (S5P) was used. Precipitated DNA was amplified by semi-quantitative PCR using primers specific for *Il6* or *Cxcl1* promoter or middle region (C) and precipitated DNA was quantified by real-time PCR using primers specific for *Il6*, *Il12b* or *Cxcl1* promoter region (D). ChIP values were normalized against the input and expressed as relative enrichment of the material precipitated by the indicated antibody on specific promoter (relative quantification using the comparative Ct method ($2^{-\Delta\Delta C_t}$)). Error bars indicate mean \pm std. deviation. The results are representative of at least three independent experiments. (E) Microarray analysis was performed in PECs from LysM-Cre⁺;Akirin2^{fl/+} and LysM-Cre⁺;Akirin2^{fl/fl} mice stimulated with LPS (100 ng/ml) for indicated time period using Affymetrix mouse genome 430 2.0 microarray chips. Robust multichip average (RMA) expression values were calculated using R package. For each probe the changes in expression between LysM-Cre⁺;Akirin2^{fl/+} and LysM-Cre⁺;Akirin2^{fl/fl} samples were defined as the difference between log₂ values for LysM-Cre⁺;Akirin2^{fl/+} and LysM-Cre⁺;Akirin2^{fl/fl} macrophages. The genes upregulated more than two fold in LysM-Cre⁺;Akirin2^{fl/+} macrophages after stimulation were selected. Hierarchical clustering classified the LPS-

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inducible genes to 8 different clusters based on the expression levels in $\text{LysM-Cre}^+; \text{Akirin2}^{\text{fl}/+}$ and $\text{LysM-Cre}^+; \text{Akirin2}^{\text{fl}/\text{fl}}$ macrophages. Statistical significance was determined using the Students *t*-test. **p* < 0.05; ***p* < 0.01.

Figure 4: Classification of LPS-induced Akirin2-dependent and independent genes

(A) 73 genes that are potentially induced by LPS in mouse peritoneal macrophages are shown. Columns 2 to 4 show the difference (log2 ratios) in expression between $\text{LysM-Cre}^+; \text{Akirin2}^{\text{fl}/+}$ and $\text{LysM-Cre}^+; \text{Akirin2}^{\text{fl}/\text{fl}}$ macrophages, for time points 0h, 2h, and 4h. Column 5 shows the cluster index assigned by hierarchical clustering (see **Figure 3E**). Columns 6, 7 and 8, 9 show the CpG scores and GC content in the regions from -100 to -1 and from +1 to 100 relative to the transcription start sites. Column 10 and 11 show whether a CpG island is predicted (1) or not (0) within the regions -10 kb to +10 kb and -1 kb to +1 kb. Colors reflect the difference in gene expression, the CpG scores, and the GC content, as indicated in the respective color legends. (B) These genes were further classified as Akirin2 dependent (lower frequency of CpG-islands) and Akirin2 independent (higher frequency of CpG-islands) genes. The pie charts show the fraction of genes associated with CpG-islands on the region -1 kb to +1 kb for Akirin2-dependent and Akirin2-independent genes, as well as the entire LPS-inducible genes and genome-wide set of genes. (C) Restriction enzyme accessibility assay was used to monitor LPS-induced nucleosome remodeling at the *Il6* promoter. PECs from $\text{LysM-Cre}^+; \text{Akirin2}^{\text{fl}/+}$ and $\text{LysM-Cre}^+; \text{Akirin2}^{\text{fl}/\text{fl}}$ were treated with LPS (100 ng/ml) for the indicated time period. Isolated nuclei were digested with *AflIII*, and the cleaved DNA were analyzed by Southern blot using *Il6* promoter specific probe. (D, E) ChIP experiments were performed with chromatin prepared from $\text{LysM-Cre}^+; \text{Akirin2}^{\text{fl}/+}$ and $\text{LysM-Cre}^+; \text{Akirin2}^{\text{fl}/\text{fl}}$ PECs treated with or without LPS (1 $\mu\text{g}/\text{ml}$) for indicated time period. Antibody against trimethyl histone ($\text{H3K4}(\text{CH}_3)_3$) (D) and acetylated histone (H3K9Ac) (E) was used. Precipitated DNA was quantified by real-time PCR using primers specific for *Il6*, *Il12b* or *Cxcl1* promoter region. ChIP values were normalized against the input and expressed as relative enrichment of the material precipitated by the indicated antibody on specific promoter (relative quantification using the comparative

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Ct method ($2^{-\Delta\Delta C_t}$). Error bars indicate mean \pm std. deviation. The results are representative of at least three independent experiments. Statistical significance was determined using the Students *t*-test. **p* < 0.05.

Figure 5: SWI/SNF components are required for the expression and histone modification of the *IL6*, but not *IL8*, promoter in response to IL-1 β stimulation.

(A) Cell lysates prepared from HEK293 cells transfected with indicated expression plasmids were immunoprecipitated with anti-FLAG antibody, followed by immunoblot analysis using anti-Myc and anti-Flag antibodies. (B) PECs from LysM-Cre⁺; *Akirin2*^{fl/+} mice were treated with LPS (1 μ g/ml) for 4 hours. Cell lysates were collected and subjected to immunoprecipitation using anti-Flag and anti-Baf60a, followed by immunoblot with anti-Akirin2 and anti-Baf60a antibodies. (C) PECs from LysM-Cre⁺; *Akirin2*^{fl/+} and LysM-Cre⁺; *Akirin2*^{fl/+} mice were treated with LPS (1 μ g/ml) for 4 hours. Cell lysates were collected and subjected to immunoprecipitation using anti-Baf60a, followed by immunoblot with anti-Akirin2 and anti-Baf60a antibodies. (D, E) HeLa cells knocked down with indicated siRNAs as shown in **Supplementary Figure S8A** were stimulated with recombinant human IL-1 β (10 ng/ml) for indicated time period. Cells were harvested for RNA isolation and cDNA was subjected to quantitative PCR analysis for gene expression of *IL6* (D) and *IL8* (E). Samples were normalized using 18S rRNA. (F, G) HeLa cells knocked down with siRNAs for indicated genes were stimulated with recombinant human IL-1 β (10 ng/ml) for indicated time period. Cells were fixed and chromatin was prepared and ChIP experiments were performed. Antibodies against phospho-RNA Pol II (S5P) (F) and Histone H3 (acetyl K9) (G) were used. Precipitated DNA was quantified by real-time PCR using primers specific for *IL6* or *IL8* promoter region. ChIP values were normalized against the input and expressed as relative enrichment of the material precipitated by the indicated antibody on specific promoter (relative quantification using the comparative Ct method ($2^{-\Delta\Delta C_t}$)). Error bars indicate mean \pm std. deviation. The results are representative of at least three independent experiments. Statistical significance was determined using the Students *t*-test. **p* < 0.05; ***p* < 0.01.

Figure 6: Akirin2 interacts with IκB-ζ via the C-terminal conserved region.

(A) Schematic representation of Akirin2 conserved regions and mutant proteins used. (B) Flag-IκB-ζ and/or Myc-Akirin2 were transfected in HeLa cells, and cell lysates were immunoprecipitated with anti-Myc Ab followed by immunoblot with anti-Flag and anti-Myc Ab. (C) HeLa cells were transfected with Flag-Akirin2 plasmid. The cells were then stimulated with IL-1β for 12 hours followed by the preparation of the cell lysates, and immunoprecipitation with anti-Flag Ab followed by immunoblot with anti-NF-κB p50, anti-NF-κB p65, anti-IκB-ζ and anti-Flag Ab. (D) PECs from *LysM-Cre⁺;Akirin2^{fl/+}* and *LysM-Cre⁺;Akirin2^{fl/fl}* mice were treated with LPS (1 μg/ml) for 4 hours. Cell lysates were collected and subjected to immunoprecipitation using anti-IκB-ζ Ab followed by immunoblot with anti-Akirin2 and anti-IκB-ζ Ab. PECs from *IκBζ^{+/+}* and *IκBζ^{-/-}* mice were harvested and cell lysates were subjected to immunoprecipitation using anti-IκB-ζ Ab followed by immunoblot with anti-Akirin2 and anti-IκB-ζ Ab. (E) Immunofluorescent staining of HeLa cells expressing Flag-Akirin2 mutants (green) together with DAPI (blue). (F) HEK293 cells were transfected with indicated Myc-tagged Akirin2 mutant plasmids and Flag-tagged IκB-ζ. Cell lysates were collected and subjected to immunoprecipitation with anti-Flag Ab followed by immunoblot with anti-Myc and anti-Flag Ab. (G) HeLa cells transfected with indicated Akirin2 mutant plasmids were stimulated with IL-1β for 12 hours followed by the preparation of the cell lysates. Then immunoprecipitation with anti-IκB-ζ Ab was performed followed by immunoblot with anti-Flag Ab. (H) HEK293 cells were transfected with indicated Flag-tagged IκB-ζ mutant plasmids and Myc-tagged Akirin2. Cell lysates were collected and subjected to immunoprecipitation with anti-Flag Ab followed by immunoblot with anti-Myc and anti-Flag Ab. (I) HEK293 cells were transfected with indicated Flag-tagged Akirin2 mutant plasmids and Myc-tagged Baf60a. Cell lysates were collected and subjected to immunoprecipitation with anti-IκB-ζ Ab followed by immunoblot with anti-Myc, anti-Flag and anti-IκB-ζ Ab. The results are representative of at least three independent experiments.

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Figure 7: The I κ B- ζ -Akirin2 cascade is critical for the recruitment of Brg1 to the *Il6* promoter for transactivation.

(A) MEFs from *Akirin2*^{fl/+} and *Akirin2*^{fl/-} mice were infected with retrovirus expressing Cre and/or full length Akirin2. Then the cells were stimulated with LPS (1 μ g/ml) followed by Northern blot analysis with indicated probes. (B, C) MEFs from *Akirin2*^{fl/-} mice were infected with retrovirus expressing Cre and indicated Akirin2 mutants. Then the cells were stimulated with LPS (1 μ g/ml) (B) and IL-1 β (10 ng/ml) (C) and the production of IL-6 was determined by ELISA. (D) ChIP experiments were performed with chromatin prepared from LysM-Cre⁺; *Akirin2*^{fl/+}, LysM-Cre⁺; *Akirin2*^{fl/fl} and I κ B- ζ ^{-/-} PECs treated with or without LPS (1 μ g/ml) for indicated time period. Antibody against Brg1 (J1) was used. Precipitated DNA was quantified by real-time PCR using primers specific for *Il6*, *Il12b* or *Cxcl1* promoter region. (E and F). Retrovirally transduced I κ B- ζ ^{-/-} (E) and *NF- κ Bp50*^{-/-} (F) BMDM expressing Flag-tagged Akirin2 were then stimulated with LPS (1 μ g/ml) for 4 hours and ChIP experiments were performed. Antibody against anti-Flag was used. Precipitated DNA was quantified by real-time PCR using primers specific for *Il6*, *Il12b* or *Cxcl1* promoter region. (G) BM Cells from LysM-Cre⁺; *Akirin2*^{fl/+} and LysM-Cre⁺; *Akirin2*^{fl/fl} mice were infected with retrovirus expressing Flag-tagged empty or full length I κ B- ζ . BMDM were then stimulated with LPS (1 μ g/ml) for 4 hours and ChIP experiments were performed. Antibody against anti-Flag was used. Precipitated DNA was quantified by real-time PCR using primers specific for *Il6*, *Il12b* or *Cxcl1* promoter region. ChIP values were normalized against the input and expressed as relative enrichment of the material precipitated by the indicated antibody on specific promoter (relative quantification using the comparative Ct method ($2^{-\Delta\Delta C_t}$)). Error bars indicate mean \pm std. deviation. The results are representative of at least two independent experiments.



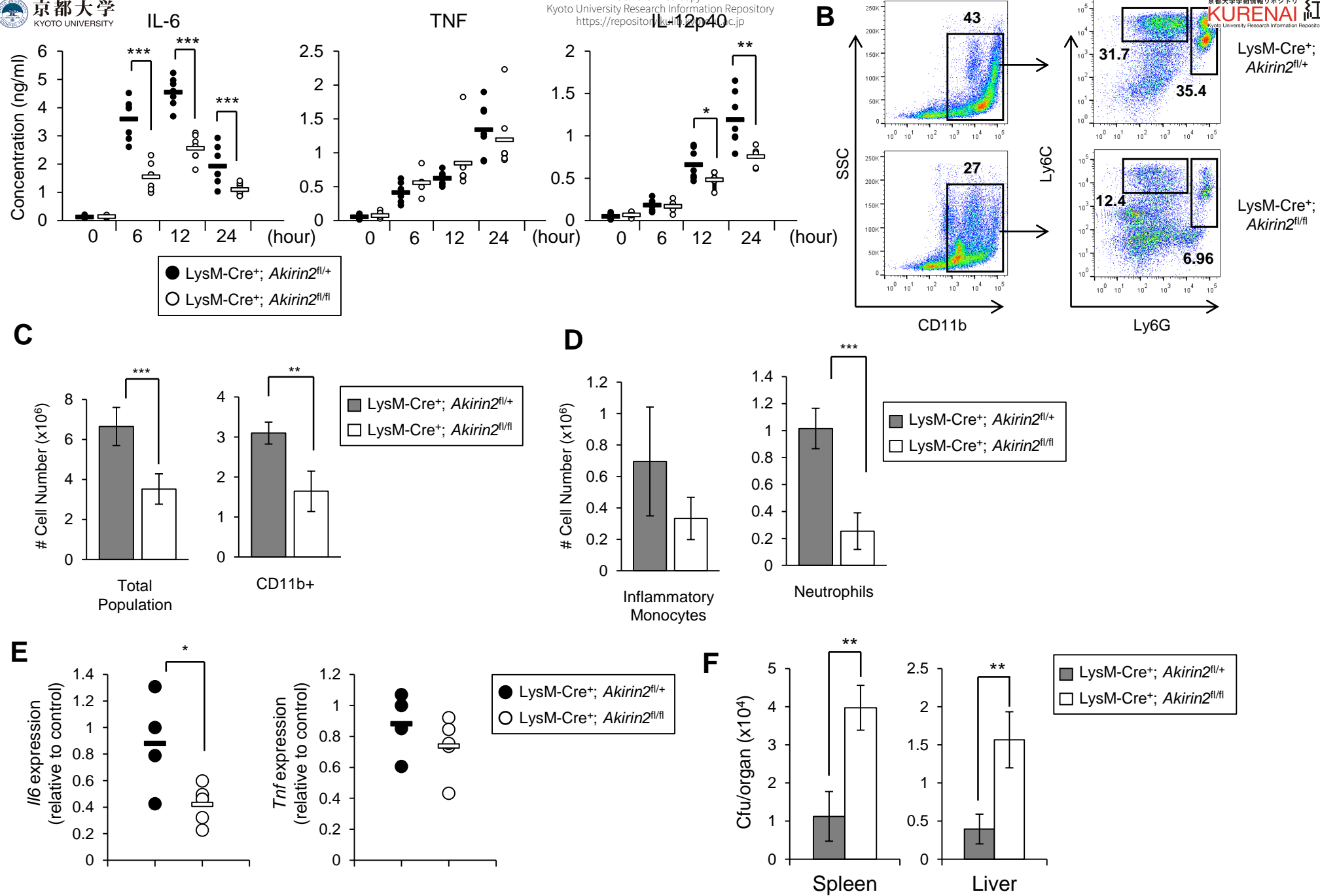


Figure 2

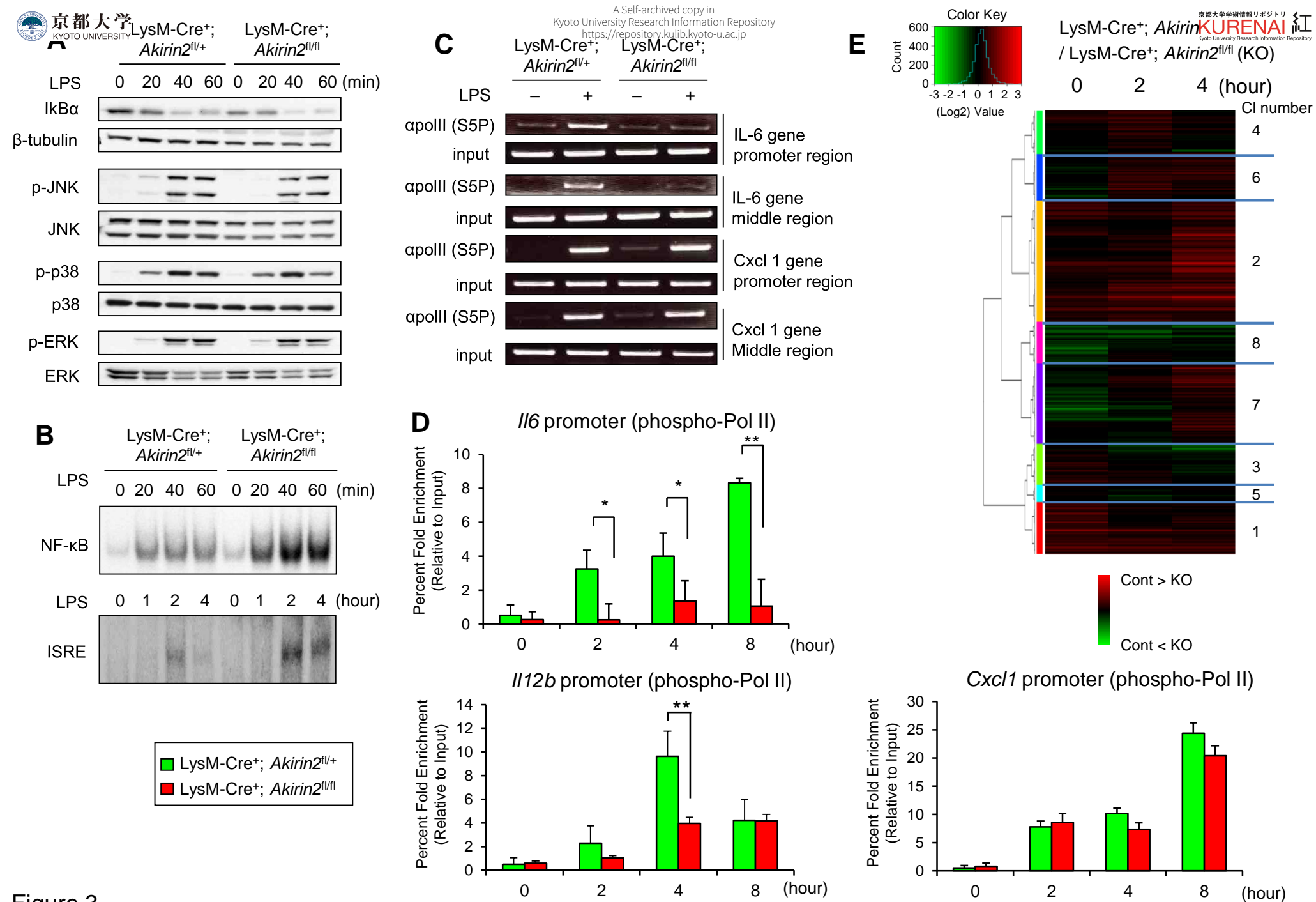
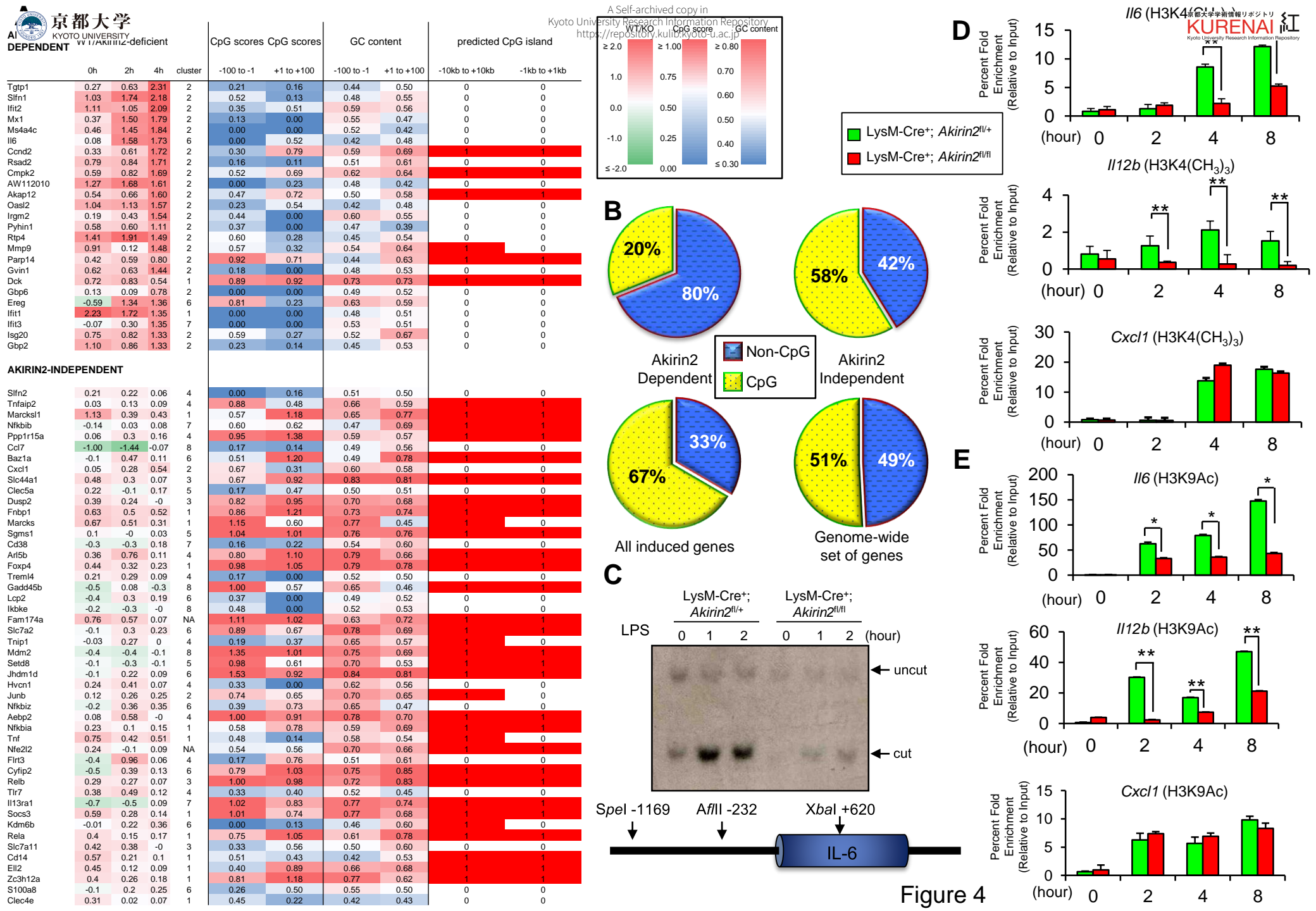
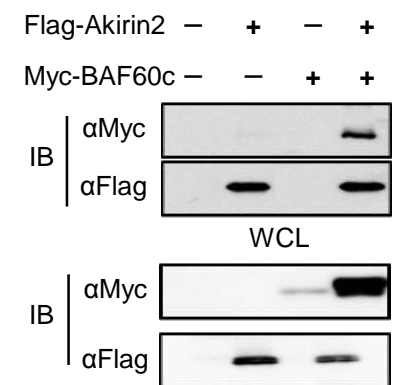
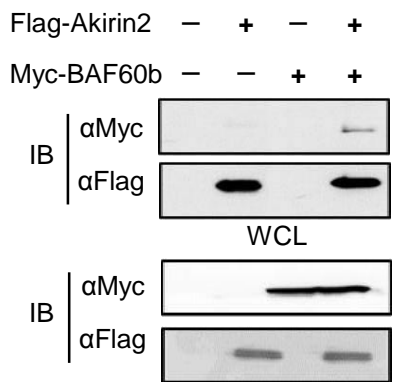
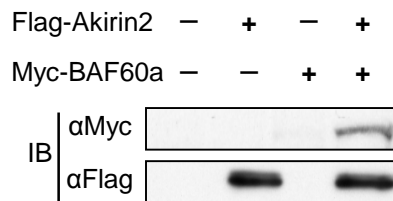
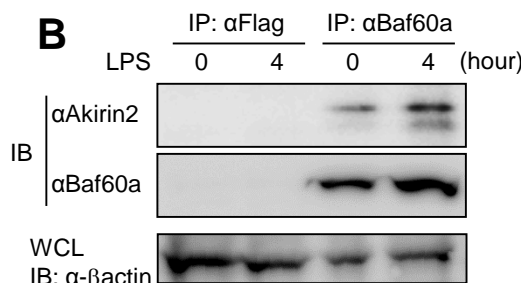


Figure 3

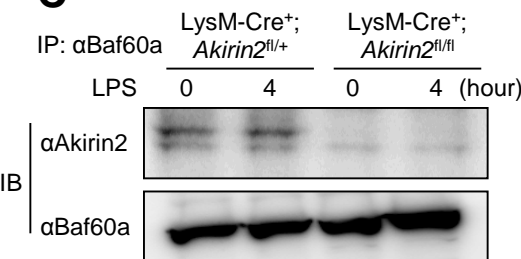




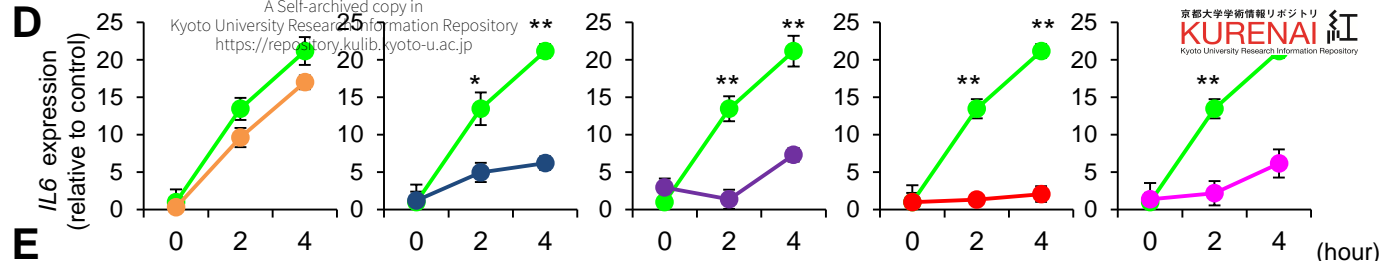
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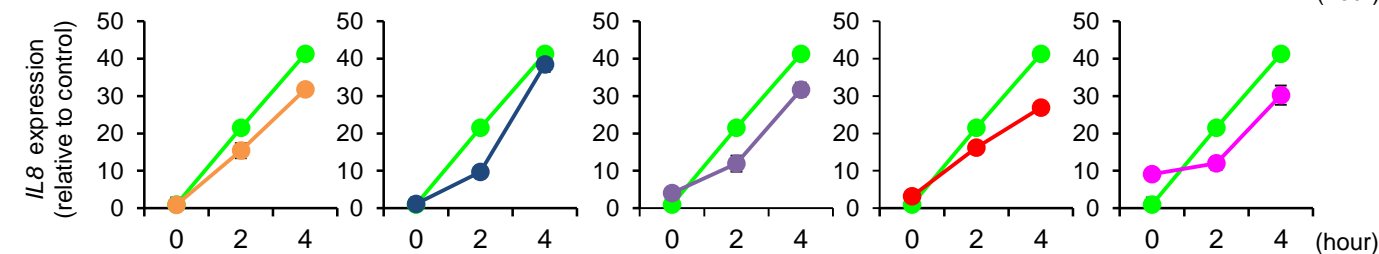
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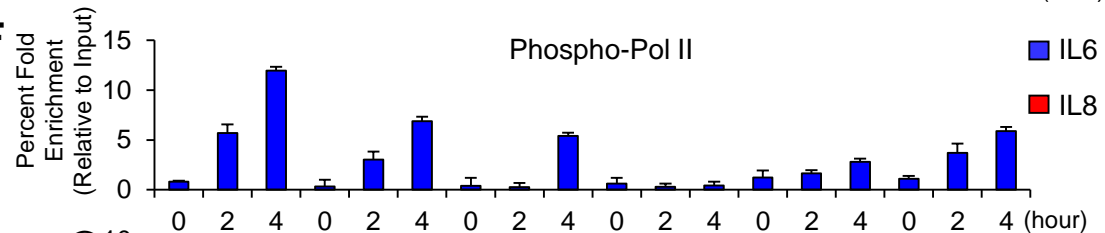
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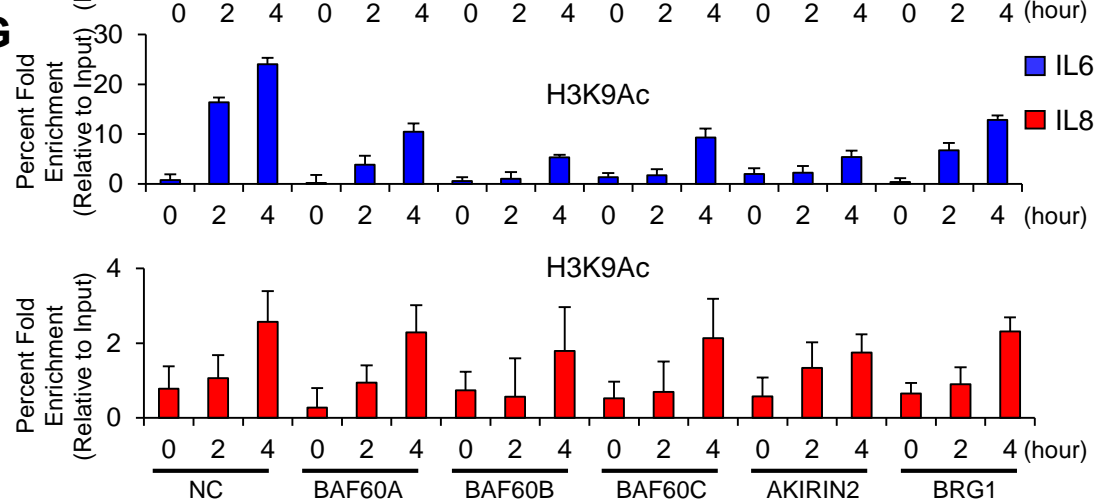


Figure 5

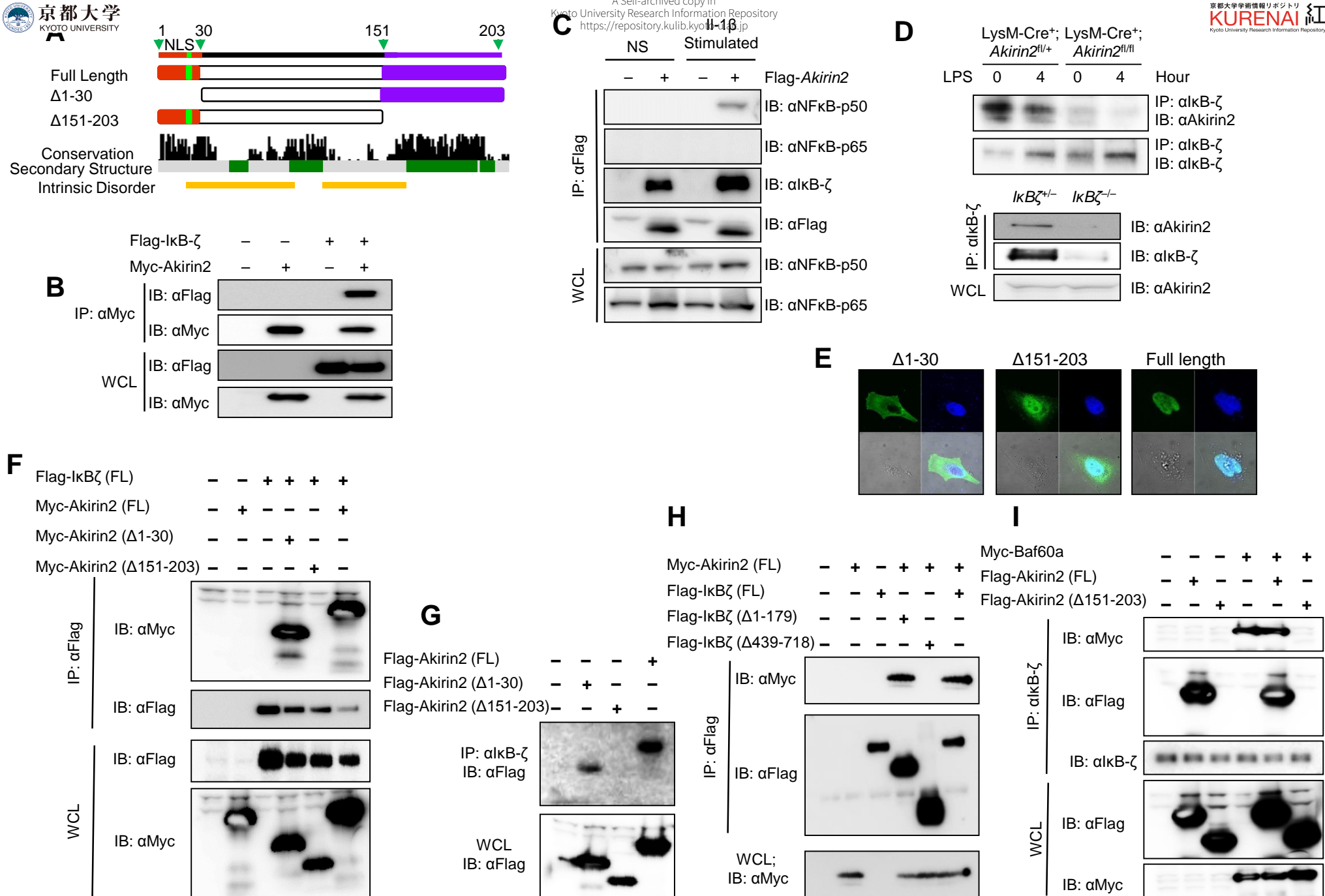


Figure 6

